AMENDMENTS TO THE CLAIMS

- 1. (Currently amended) An adjuvant comprising a <u>cationic</u> surfactant and <u>the an antigenic</u> <u>component comprising an</u> apolar fraction or part of the apolar fraction of <u>the a</u> total lipid extract of a mycobacterium, e.g. the BCG, M. microti, M. tuberculosis and M. vaccae.
- 2. (Currently amended) An adjuvant according to claim 1 where , wherein the part of the apolar fraction of the lipid extract ean be is selected from the group consisting of phthiocerol dimycocerosates, trehalose mycolipenates, glycosylated phenol phthiocerols (including phenolic glycolipids, PGL's), thehalose mycolates, sulfolipids, triacylglycerols or and menaquinones.
- 3. (Canceled)
- 4. (Currently amended) An adjuvant according to claim 3 1, wherein where the surfactant is DDA, DODA, DC-chol or DOTAP.
- 5. (Canceled)
- 6. (Currently amended) A vaccine comprising an adjuvant according to claim 1-5 1.
- 7. (Currently amended) A vaccine according to claim 6 for parenterally parenteral, oral or mucosal administration.
- 8. (Currently amended) A vaccine according to claim 7 6, wherein where the antigenic component comprises an antigenic epitope from a virulent mycobacterium, e.g. Mycobacterium tuberculosis, M. bovis or M. africanum.
- 9. (Currently amended) A vaccine according to claim 8 where , wherein the antigenic component is comprises an ESAT6-Ag85B hybrid or a fragment thereof.

- 10. (Currently amended) A An improved vaccine according to claim 7 for treating cancer, allergy or an autoimmune diseases disease, wherein the improvement comprises the adjuvant of claim 1.
- 11. (Currently amended) A delivery system comprising an adjuvant according to Claim 1–5 1.
- 12. (Currently amended) Preparing A method of preparing an adjuvant according to claim 1-5 claim 1 using thin lipid film method comprising:

dissolving a cationic surfactant and an antigenic component that comprises

an apolar fraction or part of the apolar fraction of a total lipid extract of a

mycobacterium in a solvent;

evaporating said solvent from said dissolved cationic surfactant and antigenic component with a gas;

drying said cationic surfactant and said antigenic component;

bringing said cationic surfactant and said antigenic component into a solution so as to form a thin lipid film; and

formulating the adjuvant of claim 1 from said thin lipid film.

- 13. (New) The adjuvant of claim 1, wherein said mycobacterium is BCG, M. microti, M. tuberculosis or M. vaccae.
- 14. (New) The adjuvant of claim 2, wherein said glycosylated phenol phthiocerols are phenolic glycolipids.
- 15. (New) The vaccine of claim 8, wherein said virulent bacterium is selected from the group consisting of *M. tuberculosis*, *M. bovis* and *M. africanum*.
- 16. (New) An adjuvant comprising a neutral or anionic surfactant and an antigenic component comprising an apolar fraction or part of the apolar fraction of a total lipid extract of a mycobacterium.

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17. (New) An adjuvant according to claim 16 wherein the surfactant is DOPE/PC or DOPE/PC/PG.